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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 2 of 2

Complete if Known	
Application Number	09/308,080
Filing Date	October 28, 1999
First Named Inventor	Frank Gonzalez
Group Art Unit	1652
Examiner Name	Steadman, D
Attorney Docket Number	015280-271100US

OTHER PRIOR ART -- NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
DJS	AC	Gonzalez, F., et al., "Diagnostic analysis, clinical importance and molecular basis of dihydropyrimidine dehydrogenase deficiency," TiPS, 16:325-327 (1995).	
	AD	Hiroshi Y., et al. "cDNA cloning and chromosome mapping of human dihydropyrimidine dehydrogenase, an enzyme associated with 5-flourouracil toxicity and congenital thymine uraciluria," J. Biol. Chem., 269:23192-23196 (1994).	
	AE	JOURNAL BIOLOGICAL CHEMISTRY, vol. 264, no. 20, July 1990, pages 12067-74, XP002032866 KUIVANIEMI, H., ET AL: "Identical G to a mutations in three different introns of the type III procollagen gene (COL3A1) produce different patterns of RNA splicing in three variants of Ehlers-Danlos Syndrome IV," see abstract	
	AF	Meinsma, R., et al., "Human Polymorphism in Drug Metabolism: Mutation in the Dihydropyrimidine Dehydrogenase Gene Resutls in Exon Skipping and Thymine Uracilurea," DNA & Cell. Biol., 14(1)1-6 (1995).	
	AG	NUCLEIC ACIDS RESEARCH, vol. 15 no. 14, 1987, pages 5613-28, XP002032865 MARVIT, J. ET AL: "GT to AT transition at a splice donor site causes skipping of the preceeding exon in penylketonuria" see abstract	
	AH	SINGAPORE JOURNAL OF OBSTETRICS AND GYNECOLOGY, vol. 26, no. 3, November 1995, pages 176-86, XP000600337 ROY ET AL: "molecular scanning of human diseases" see the whole document.	
	AI	Vreken, P., et al., "A point mutation in an invariant splice donor site leads to exon skipping in two unrelated Dutch patients with dihydropyrimidine dehydrogenase deficiency," J. Inherit. Metab. Dis., 19(5):645-54 (1996).	
DJS	AJ	Wei, X., et al. "Molecular Basis of the Human Dihydropyrimidine Dehydrogenase Deficiency and 5-Fluorouracil Toxicity," J. Clin. Invest., 98(3)610-615 (1996).	

Examiner Signature	DJS	Date Considered	6/30/02

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[1] Unique citation designation number. [2] Applicant is to place a check mark here if English language Translation is attached.

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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Sheet 1 of 2

	Complete if Known
Application Number	09/308,080
Filing Date	October 28, 1999
First Named Inventor	Frank Gonzalez
Group Art Unit	1652
Examiner Name	Steadman, D
Attorney Docket Number	015280-271100US

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	U.S. Patent Document Number	Kind Code[2] (*if known*)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
DJS	AA	5,856,454		Gonzalez et al.	01/05/1999	536/23.1

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document Office[3]	Number[4]	Kind Code[5] (*if known*)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]
DJS	AB		WO 97/35034		Gonzalez et al.	09/25/1997		

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Examiner Signature	DJS	Date Considered	6/30/02

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